

Q1
AMB
(c) causes a clinically significant improvement in a standard marker of renal function
when administered to a mammal in, or at risk of, chronic renal failure.

13. A method as in [any one of claims 1-12] claim 1 wherein
said mammal is afflicted with a condition selected from the group consisting of chronic
renal failure, end-stage renal disease, chronic diabetic nephropathy, diabetic glomerulopathy,
diabetic renal hypertrophy, hypertensive nephrosclerosis, hypertensive glomerulosclerosis,
chronic glomerulonephritis, hereditary nephritis, and renal dysplasia.

Q2
14. A method as in [any one of claims 1-12] claim 1 wherein
examination of a renal biopsy of said mammal indicates that said mammal is afflicted
with a condition selected from the group consisting of glomerular hypertrophy, tubular
hypertrophy, glomerulosclerosis, and tubulointerstitial sclerosis.

15. A method as in [any one of claims 1-12] claim 1 wherein
examination of said mammal indicates renal fibrosis.

July 17
Q3
17. A method as in [any one of claims 1-12] claim 1 wherein
said mammal possesses a number of functional nephron units which is less than about
50% of a number of functional nephron units present in a mammal having intact healthy
kidneys.

Q4
24. A method as in [any one of claims 1-12] claim 1 wherein
said mammal has a GFR which is chronically less than about 50% of a GFR_{exp} for said
mammal.

Q5
28. A method as in [any one of claims 1-12] claim 1 wherein
said mammal is a human male weighing at least about 50 kg and has a GFR which is
chronically less than about 50 ml/min.